

NASA CR-54782 MRB5010Q1

Quarterly Report No. 1

RESEARCH AND DEVELOPMENT OF THE DRY TAPE BATTERY CONCEPT

9 June to 8 September 1965

N 66 - 12 16	9 (THRW)	CFSTI PRICE(S) \$
(PAGES)	(CODE)	Hard copy (HC) 3.00
(NASA CR OR TMX OR AD NUMBER)	(CATEGORY)	Microfiche (MF) 50
	by	ff 653 July 65

J. J. Byrne, L. F. Athearn, J. C. Mac Donald and D. L. Williams

prepared for

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

CONTRACT NAS3-7624

MONSANTO RESEARCH CORPORATION
BOSTON LABORATORY
Everett, Massachusetts 02149
Tel: 617-389-0480

Quarterly Report No. 1
"RESEARCH AND DEVELOPMENT OF THE DRY TAPE BATTERY CONCEPT"
9 June to 8 September 1965

Contract No. NAS3-7624 8 October 1965

MRC Report No. MRB5010Q1

For

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION Lewis Research Center Cleveland, Ohio

Authors

Joseph J. Byrne, Project Leader Lee F. Athearn John C. Mac Donald David L. Williams

Technical Management
NASA Lewis Research Center
Cleveland, Ohio
Space Power Division
W. J. Nagle, Technical Manager

MONSANTO RESEARCH CORPORATION BOSTON LABORATORY Everett, Massachusetts 02149 Tel: 617-389-0480

SUMMARY

A new tape couple, based on the Mg/AlCl₃,HCl/trichloro-triazinetrione (Monsanto ACL-85³) system, has been developed. This cell discharges at 2.6 volts at 0.2A/in.² and 2.2 volts at 0.8 A/in.². Cathode coulombic efficiencies range from 60-80%, with individual test results as high as 87%. The discharge curve is flat and current density has little effect on cathode efficiency. This tape couple can tolerate current drains in excess of lA/in.², but cell voltage falls below 2.0 volts.

The cathode tape configuration is identical to that developed in our previous program (NAS3-4168) except that less inert material is required with ACL-85 compared with previous (potassium periodate, picric acid) cathode materials. Present cathodes operate satisfactorily with up to 75 wt-% active material. Maximum active material content in previous systems was 60 wt-%.

Energy output from the Mg/ACL-85 system is substantially higher than any other system studied to date. Accurate electrolyte requirement data have not yet been determined but, on a dry basis, delivered energy densities over 300 watt-hour/pound of complete tape cell, excluding electrolyte, have been obtained.

A satisfactory method was developed for preparing ACL-85 tape cathodes from a trichloroethylene slurry. Vacuum pretreatment of the acetylene black conductor is required to prevent hydrolytic decomposition of the active chlorine material during tape processing and storage.

The best nonaqueous electrolytes for lithium were found to be potassium hexafluorophosphate and lithium perchlorate in methyl formate. Anode half cell voltages of -2.1 to -2.3 volts vs. NCE could be maintained for extended periods at 0.5 amp/in.². Results for magnesium in nonaqueous electrolytes were erratic due primarily to sensitivity to anode surface preparation.

Vapor phase chromatographic techniques for water analysis are being developed. It appears that concentrations as low as 10 ppm can be determined routinely for most electrolyte systems.

New electroanalytical test equipment has been designed and built for both static and dynamic tests. Fast switching pulse current equipment is now available to record the trace of electrode performance during the first ten milliseconds of discharge. In addition, IR-free voltage measurements can be made at currents up to three amperes. A new dynamic test station was built that is capable of automatic data recording, tape speed control, and electrolyte input. The unit can run unattended for 24 hours.

TABLE OF CONTENTS

			Page
I.	INI	TRODUCTION	1
	A. B. C.	CONTRACT OBJECTIVES	1 1 2
II.	TAS	K I. HIGH ENERGY COUPLE RESEARCH	3
	Α.	CATHODE RESEARCH AND DEVELOPMENT	3
		1. 2,4,6-Tri-N-Chlorotriazinetrione in Aqueous Electrolyte	3
		2. 2,4,6-Tri-N-Chlorotriazinetrione in Nonaqueous Electrolyte	10
		3. Plans for Second Quarter	14
	B.	ANODE RESEARCH AND DEVELOPMENT	14
		l. Lithium in Nonaqueous Electrolyte	14
		2. Magnesium in Nonaqueous Electrolyte	21
		3. Plans for Second Quarter	23
III.	TAS	K II. TAPE CELL EVALUATION	24
	A. B.		24 24
IV.	TAS	K III. SUPPORTING RESEARCH	25
	Α.	TEST EQUIPMENT DESIGN AND IMPROVEMENT	25
		l. Free Electrolyte Half Cell Unit	25
		2. Limited Electrolyte Full Cell Static Test Unit.	25
	•	3. Dynamic Test Station	25
		4. Improved Water Analysis Techniques	· 31
٧.	REF	PERENCES	40
	ДРР	PENDTX	41

LIST OF FIGURES

No.		Page
1.	Effect of Electrolyte on Polarization of 2,4,6- Trichlorotriazinetrione Cathode	6
2.	Effect of Active Material Concentration in Cathode Mix on Polarization.	7
3.	Discharge Characteristics of Mg/Trichlorotriazine-trione Tape Cell in Various Electrolytes.	8
4.	Discharge Characteristics of Mg/Trichlorotriazine-trione Tape Cells in Mixed Electrolytes.	9
5.	Discharge Characteristics of ACL- $85^{\$}$ Tape Cells at Various Current Densities.	12
6.	Oscilloscope Traces of Lithium and Magnesium in $\gamma\textsc{-Butyrolactone}$ (lM KPFs) Electrolyte.	16
7.	Polarization Curves of Lithium in Methyl Formate.	20
8.	Circuitry for Free Electrolyte Half Cell Test.	26
9.	Free Electrolyte Test Cell.	27
10.	Kordesch-Marko Type Interrupter Circuit Diagram.	28
11.	Allis-Chalmers Type Interrupter Circuit Diagram.	29
12.	Dynamic Test Unit Drive, Recorder and Electrolyte Supply.	30
13.	Dimethyl Formamide Vapor Phase Chromatogram.	33
14.	Methanol Chromatogram.	34
15.	Methyl Formate Chromatogram.	35
16.	Ethyl Formate Chromatogram.	36
17.	Y-Butyrolactone Chromatogram.	37
18.	Acetonitrile Chromatogram.	38

LIST OF TABLES

No.		Page
1.	Discharge Characteristics of Potassium Periodate Cathode Tapes Prepared from Trichloroethylene Slurries and Vacuum Dried.	4
2.	The Effect of Acetylene Black Pretreatment on ACL-85 Stability.	5
3.	Typical Discharge Data for Mg/AlCl3, HC1/ACL-85 Systems.	11
4.	Energy Density Calculation from Static Cell Discharge Data.	13
5.	Solubility and Observable Stability of ACL-85 in Various Organic Solvents.	14
6.	Free Electrolyte Cell Discharge Data for Lithium in Nonaqueous Electrolyte.	17
7.	Free Electrolyte Cell Discharge Data for Magnesium in Nonaqueous Electrolyte.	18
8.	Current-Voltage Data for the Mg/Mg(ClO ₄) ₂ , Methanol System.	23
9.	Preliminary Calibration Data for Vapor Phase Chromatographic Determination of Water.	39
10.	Solvent Retention Times.	39
	APPENDIX TABLES	
A-l	Static Cell Discharge for Tri-N-Chlorotriazine-trione Tape Cathodes.	41
A-2	Analytical Method for the Determination of Available Chlorine.	45

I. <u>INTRODUCTION</u>

A. BACKGROUND

This work is a continuation of research performed under Contracts NAS3-2777 and NAS3-4168. During the initial program (NAS3-2777) the feasibility of the Dry Tape Concept was demonstrated using a divalent silver oxide-coated, porous polypropylene tape that was drawn between two current collectors. One current collector was a zinc block that also served as the anode. The cathode current collector was a thin silver plate. Electrolyte was supplied by a second tape prewetted with electrolyte and stored separately ("dual tape system"). The system was activated by bringing the two tapes together just prior to their entry into the current collector zone.

The tapes were driven by a spring-wound motor, contained in a separate housing. Within this housing was an output metering and load section. Four such drive housings, together with 20 tape decks, were supplied to NASA for demonstration. In continuous operation, silver peroxide utilizations of over 90% were obtained at current densities of 1 amp/in.².

In the follow-on program (NAS3-4168), the "dual tape system" using the established battery couple, A20/KOH/Zn, was replaced by a single tape configuration using a thin foil magnesium anode and cathode coatings containing nonconducting, higher energy organic (dinitrobenzene, picric acid, trichloro-triazinetrione) and inorganic (potassium meta-periodate) depolarizers. Efficient, high drain rate discharge of these cathode materials was achieved through use of the "thin plate tape electrode" configuration. The single tape configuration was optimized for the system, Mg/ 2MA1Cl₃;0.5M HCl/KIO₄. Up to 80% KIO₄ utilization was achieved with a cell voltage of 2.2 volts at a current density of 0.5 amp/in.2 while in a moving configuration. In addition, to tape electrode configuration development, methods of electrolyte incapsulation and tape activation were devised. Also, techniques for supplying multiple cell voltage, parasitic drive and continuous coated tape manufacture were developed.

B. <u>CONTRACT OBJECTIVES</u>

Specific objectives of this program are to:

- 1. Develop a high energy tape couple that will deliver 200 watt-hours per pound in a practical configuration.
- 2. Incorporate this couple in a tape configuration that will exhibit prescribed flexibility and abrasion resistance during operation.

The major objective is to improve substantially the energy density output of the dry tape system. Despite good electrode efficiencies, the best delivered energy densities obtained with organic nitro compound or periodate cathode systems were under 100 watt-hours per pound. This was primarily due to high electrolyte consumption and a high cathode inert weight requirement.

In the present work, only couples with extremely high potential energy densities are being considered. Couples with theoretical energy densities greater than 750 watt-hours per pound are to be incorporated into the tape electrode configurations developed in previous programs.

C. FIRST QUARTER OBJECTIVES

Objectives for this quarter were to:

- 1. Develop and evaluate a new aqueous system, based on the magnesium/2,4,6-trichlorotriazinetrione (Monsanto AC1-85) couple.
- Screen discharge characteristics of lithium and magnesium in selected organic solvent-electrolyte combinations.
 - a. Build improved half-cell testing stations.
 - b. Purify and analyze solvents and electrolyte salts.
 - c. Determine the effect of purity on conductivity.
 - d. Determine polarization and efficiency characteristics of lithium and magnesium in selected electrolyte systems.
- 3. Initiate study of stability and solubility of cathode materials in selected organic solvents.
- 4. Design, build, and test an improved dynamic test station.

II. TASK I. HIGH ENERGY COUPLE RESEARCH

A. CATHODE RESEARCH AND DEVELOPMENT

1. 2,4,6-Trichlorotriazinetrione in Aqueous Electrolyte

The theoretical energy density of the magnesium-2,4,6-tri-chlorotriazinetrione couple is estimated to be 785 watt-hours per pound of reactant, assuming the following reaction:

Previous work in this laboratory showed that this material would exhibit flat, efficient discharge in aqueous electrolyte. Preliminary work in the previous tape program (NAS3-4168) indicated that a trichlorotriazinetrione tape cathode could tolerate higher drain rates with less inert weight then the organic nitro compound and periodate based tapes.

The development program during this period was designed to optimize energy output at the highest current drains possible. Tape cell evaluations were performed using the simple static test described in our previous program. A complete tabulation of results is shown in Table A-1.

a. Tape Cathode Preparation

To avoid possible activity loss due to hydrolysis prior to discharge a method for cathode tape preparation was developed by which the cathode mix is cast or extruded onto the tape from a trichloroethylene slurry and dried under vacuum at 60-70°C. The acetylene black, active cathode material (Monsanto ACL-85) and reinforcing fibers (carbon or dynel) are dry-blended by stirring. The acetylene black and carbon fibers are prefluffed in a Waring Blendor prior to use. The polyvinyl formal binder is added as a trichloroethylene solution (0.005 g PVF/ml solution). The resulting slurry is thoroughly mixed by mechanical stirring and hand cast onto the separator tape, using a doctor blade technique. The same slurry may be used for continuous tape manufacture by the extrusion process developed in the previous contract. Hand-cast tapes are vacuum-dried for one hour at 60-70°C.

This technique initially was tested using the potassium periodate cathode formulation developed in our prior program (NAS3-

4168). Discharge results summarized in Table 1 show somewhat improved output and reproducibility compared with similar periodate tapes made from adueous slurries and then air dried.

Table 1

DISCHARGE CHARACTERISTICS OF POTASSIUM PERIODATE CATHODE TAPES PREPARED FROM TRICHLOROETHYLENE SLURRIES AND VACUUM DRIED

54 wt-% KIO₄, 40.5 wt-% acetylene black, (50% compressed) 2.75 wt-% polyvinyl formal, 2 75 wt-% carbon fibers, 3.5 mil dynel tape.

primary magnesi; um foil anode - 2M AlCl3 · 0.5M HCl electrolyte

	Operating Cell Voltage, volts	Cathode Effici- ency, %	Current Density, amp/in.2
Summary of eight tests	2.14 <u>+</u> 0.07	70 <u>+</u> 7%	0.5

This procedure works well for ACL-85. However, it was found that a vacuum drying pretreatment (12 hours, 200°C) of the acetylene black is required to prevent substantial cathode activity loss during processing and storage. The reasons for this are prepresented in the following section.

b. Stability of Trichlorotriazinetrione Tape Cathode

The stability of ACL-85 tape cathode was monitored by both discharge-efficiency tests and active chlorine chemical analysis. The analytical method for determining the active chlorine content of tape cathodes consists of an iodometric titration with sodium thiosulfate to a starch endpoint. The detailed analysis is described in Table A-2.

Tapes prepared with "as received" acetylene black showed substantial activity loss during processing and on subsequent standings. It appears that water, adsorbed on Shawinigan acetylene black, is the cause of activity loss of 2,4,6-trichlorotriazinetrione (Monsanto Company ACL-85). Although ACL-85 is fairly stable to moisture in the air, the acetylene black appears to catalyze hydrolysis by adsorbed water. Vacuum drying of the acetylene black eliminated significant activity loss that previously occurred during tape processing. In addition, tapes prepared with dried acetylene black, retained this stability on standing in air at least up to seven days. Activity loss data are reported in Table 2.

Table 2.

THE EFFECT OF ACETYLENE BLACK PRETREATMENT ON ACL-85® STABILITY

	Measured Active Ch.	
	As Received	Dried Acetylene
Material and Condition	<u>Acetylene Black</u>	Black
ACL-85, as received, theoretica ACL-85, as received, analyzed		0.1
Tape (50% ACL-85), freshly prepa	red 77.1 + 0.4	85.4 ± 1.3
Tape (50% ACL-85), 24 hours in		84.0 ± 0.9
Tape (50% ACL-85), 96 hours in		82.5 \pm 1.1
Tape (50% ACL-85), 168 hours in	air 61.2 \pm 1.8	83.3 + 0.4

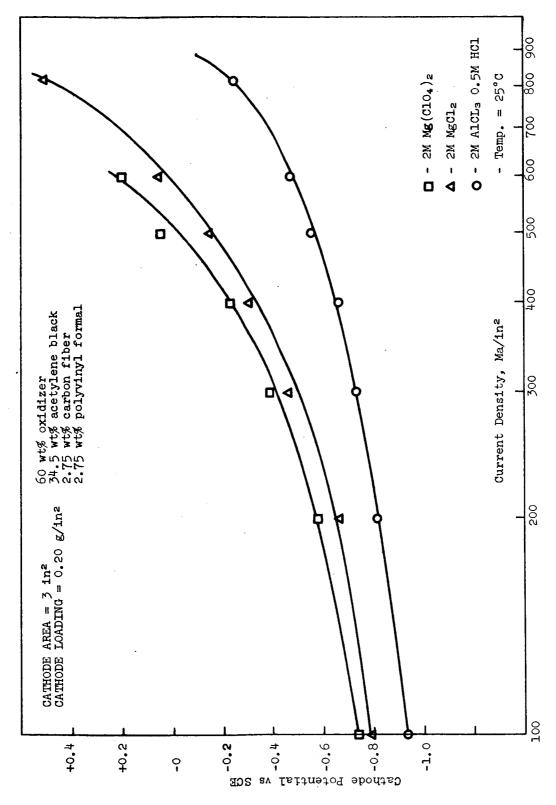
There appears to be 3-5% activity loss on processing, and a slight loss appears to occur with time during air storage, but measured losses are almost within experimental error.

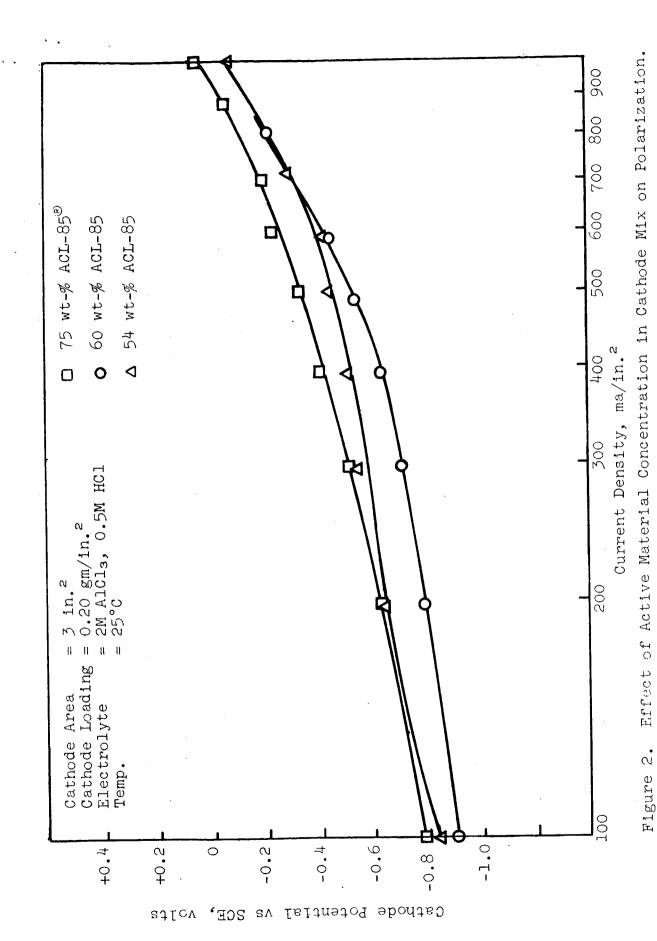
c. Discharge of ACL-85 Tape Cathodes in Aqueous Electrolyte

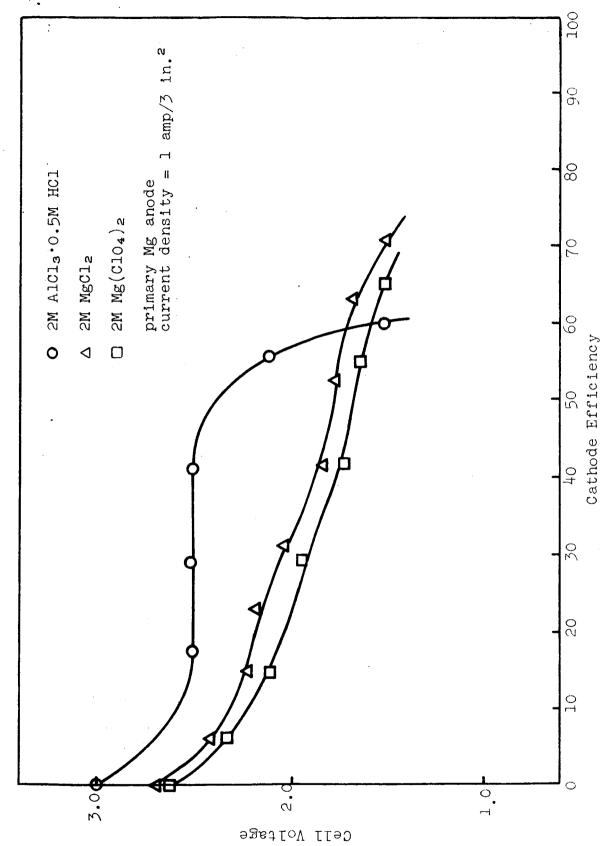
Polarization and efficiency measurements were carried out in magnesium chloride, magnesium perchlorate, and aluminum chloride-hydrochloric acid electrolyte with primary magnesium as the anode electrode. Cathode formulations varied from 50 to 80 wt-% ACL-85. Acetylene black, polyvinyl formal binder and dynel fiber comprised the remainder.

As a general rule, polarization was less severe in the aluminum chloride-hydrochloric acid electrolyte. Figure 1 is a typical illustration. There was little difference in polarization characteristics between the 50 wt-% and 75 wt-% active material concentration, as shown in Figure 2.

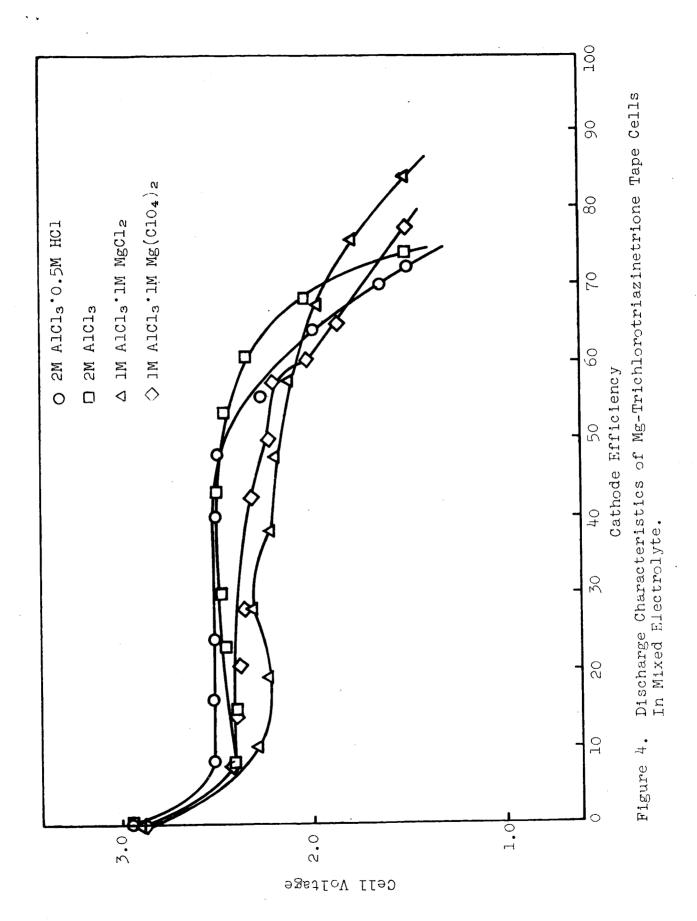
The effect of electrolyte on the shape of the discharge curve is significant. In neutral magnesium chloride or magnesium perchlorate, the voltage drops steadily. In strong acid electrolyte, a fairly constant voltage is maintained until the cell is nearly depleted. This behavior is illustrated in Figure 3. The initial emf difference is primarily a pH effect. However, the steady voltage drop during discharge is due solely to cathode polarization according to half cell measurements. The use of mixed salt electrolyte eliminates this deterioration. Figure 4 shows the effect of adding aluminum chloride to magnesium chloride and magnesium perchlorate. The initial potential dip occurring in some cases is due to the finite time required to break down the oxide layer on the anode surface.







Discharge Characteristics of Mg-Trichloro-Triazine-Trione Tape Cell in Various Electrolytes. Figure 3.



Morehouse and Glicksman (ref. 1) previously observed this phenomenon and ascribed it to poor cathode stability in neutral electrolyte. However, the results here show no significant capacity loss in neutral electrolyte. Rather, the gradual emf deterioration seems to indicate polarization caused by accumulation of some reaction product. The nature of this behavior is not known at present, and is the subject of current study.

To date, the best cathode coulombic efficiencies (70-80%) have been obtained with 65-70 wt-% ACL-85 cathode tapes. Discharge voltages vs primary magnesium range from 2.65 v at 0.170 amp/in.² to 2.20 v at 0.800 amp/in.². There is no effect on efficiency over this current density range. Some typical static cell discharge data are shown in Table 3, and are illustrated in Figure 5. All efficiency data are based on actual active chlorine content, as determined by analysis.

Accurate power density data for tape cells require precise electrolyte consumption data, which are available only from dynamic testing. In static tests, energy densities of approximately 300 watt-hours/lb of dry cell (excluding electrolyte weight) are obtained. A typical example is shown in Table 4. The delivered energy density per unit cell weight will be considerably lower when accurate electrolyte requirement data are included. These figures are useful, however, in evaluating changes in formulations and discharge conditions while still in the static cell test phase. Dynamic testing of promising ACL-85 tape cells will be initiated early in the second quarter.

2. 2,4,6-Trichlorotriazinetrione in Nonaqueous Electrolyte

Independent work in this laboratory indicated that ACL-85 was a promising cathode material for nonaqueous systems. A Li/IM LiClO₄, butyrolactone/ACL-85 cell in a tape configuration discharged at 3.1 volts under a current density of 0.07 amp/in.². During this quarter, some preliminary compatibility tests were performed prior to full-scale cell testing, which will be carried out during the second quarter.

a. Solubility of ACL-85 in Organic Solvents

Simple screening tests based on visual observation were performed to check stability of ACL-85 in several potential organic electrolyte solvents. In addition, solubilities at 25°C were determined.

Table 3

TYPICAL DISCHARGE DATA FOR Mg/AlCl₃, HCl/ACL-85[®] SYSTEMS

Cell Area = 3 in. x 1 in., Electrolyte = 2M AlCl₃ 0.5M HCl

	_	, ,		
Cell No.	ACL-85, _wt-%	Current, I, amps	Average Voltage, volts	Cathode Coulombic Efficiency, % (1.5 volt cutoff)
78673-8	65	0.5	2.65	71
78673-6	65	1.0	2.48	76
78673-1	65	1.0	2.50	75
78673-10	65	1.5	2.45	83
78673-2	65	1.5	2.38	77
78673-9	65	2.0	2.40	82
78673-3	65	2.0	2.31	84
78673-7	65	2.5	2.20	76
78677-2	70	1.0	2.50	79
78677-7	70	1.5	2.40	80
78677-6	70	1.5	2.40	80
78677-9	70	2.0	2.30	79
78677-3	70	2.0	2.25	80
78677-4	70	2.5	2.20	79
78677-1	70	3.0	1.90	68

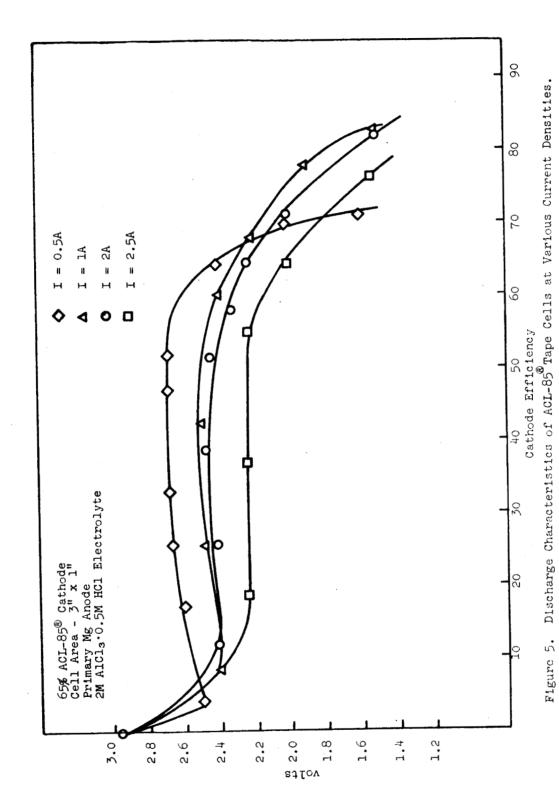


Table 4

ENERGY DENSITY CALCULATION FROM STATIC CELL DISCHARGE DATA

Cell No. 78677-6

Latinity Doct

70 wt-% ACL-85® Cathode Formulation:

24.5 wt-% Shawinigan Acetylene Black

2.75 wt-% 1/8 in. Dynel Fiber 2.75 wt-% Polyvinyl formal binder

.035 in. Dynel Separator

5-mil perforated primary Magnesium Anode:

0.9324 g Cell Weight: Cathode Coat

Separator 0.0883 g 0.1201 g Magnesium 1.1408 g

Electrolyte 2M AlCl₃ 0.5M HCl Current Density 0.5 amp/in.²

Current

= 1.5 amp (cell area = 3 in.²) = -2.92 v (cathode = 1.02 vs S.C.E.) OCV

Average Operating Voltage = 2.40 v

Power Density = $\frac{2.40 \text{ v x l.5a x l2.2min x 454 g/lb}}{}$ 1.1408 g x 60 min/hr

> = 292 watt hours/lb dry cell (excluding electrolyte)

Table 5.

SOLUBILITY AND OBSERVABLE STABILITY OF ACL-85
IN VARIOUS ORGANIC SOLVENTS

Solvent	Observed Reaction	Solubility g/100 cc at 25°C
Methanol	none	21.4
Acetone	none	10.1
Butyrolactone	none	18.2
Acetonitrile	none	11.2
Dimethyl formamide	none	46.2
Methyl formate	none	2.4

3. Plans for Second Quarter (Cathode Research and Development)

The energy density output by the aqueous Mg/ACL-85 system will be optimized and the complete tape system will be evaluated under dynamic conditions.

The performance of active chlorine compounds in nonaqueous electrolyte will be investigated and full cell tape configuration studies using a lithium foil anode will be started.

B. ANODE RESEARCH AND DEVELOPMENT

1. Lithium in Nonaqueous Electrolyte

Work during this quarter was devoted primarily to setting up suitable electrochemical test apparatus, materials purification and analysis, and screening of selected lithium-electrolyte systems.

a. Half Cell Test Apparatus for Nonaqueous Systems

In past work with aqueous systems, no attempt was made to identify quantitatively the separate components that contributed to polarization. In dealing with nonaqueous systems, IR polarization becomes important. The electrical apparatus for half cell and full cell static testing was designed to give as much quantitative information as possible about the various polarization effects.

Electrodes are evaluated initially using a free electrolyte half cell test. Constant current is used and an oscilloscope trace shows IR polarization and, in some cases, isolates concentra-

tion polarization and passivation effects. The test cell and associated circuitry are described completely in Section IV. Using this system, IR can be determined within 1 millisecond of application of current. A platinum counter electrode is employed to eliminate interfering cathode effects. In addition, the test cell is designed to measure gassing rate. Two typical oscilloscope traces are shown in Figure 6.

Nonaqueous systems in a tape configuration are evaluated using the conventional sandwich cell static test unit. To give an indication of IR polarization, interruptor-type bridges have been modified for battery work. These circuits are described in Section IV. The cell itself is constructed from polypropylene for stability in nonaqueous solvents. Working counter electrodes are silver chloride, platinum screen or in some cases ACL-85 cathode tapes. The silver chloride cathode was prepared by anodizing a discharged silver oxide electrode (Yardney Electric Silcad® YS10) in hydrochloric acid. This cathode operates satisfactorily but suffers from fragility.

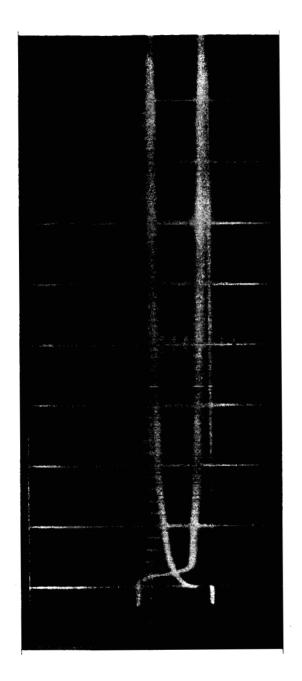
b. <u>Materials</u>

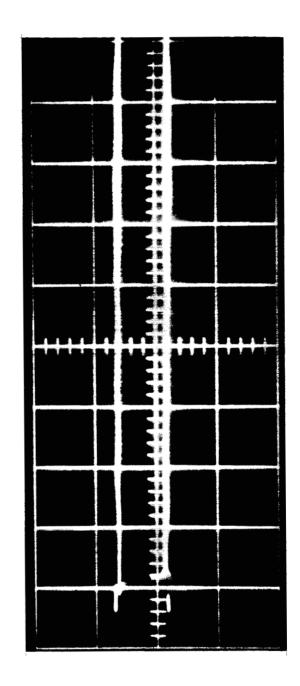
The electrolyte systems were selected on the basis of previous NASA-sponsored nonaqueous battery system research. All solvents were dried over Linde 4A Molecular Sieves. Methanol was predried over Linde 5A Molecular Sieves. The solvents were separated by vacuum distillation. Salts, except for morpholinium hexafluorophosphate, were vacuum-dried at 150°C for a minimum of 24 hours. Hygroscopic salts such as perchlorates required drying times up to 72 hours. Morpholinium hexafluorophosphate was dried in a vacuum dessicator over phosphorus pentoxide to avoid decomposition at elevated temperatures.

Dried materials transfer and electrochemical measurements were carried out in a Kewaunee Scientific Equipment (20380) drybox under an argon atmosphere. Phosphorus pentoxide drying agent is used to trap any water vapor.

Water analyses reported here were performed using a Karl Fisher amperammetric titration with a Beckman KF-3 Aquameter. A vapor phase chromatographic technique is being developed that promises to be more reproducible. Details of this method are reported in Section IV. Water concentrations below 0.5 mg $\rm H_2O/ml$ of solution, as measured by the Karl Fisher method, are questionable because of the high concentration reagents used. All water analyses are listed in Tables 6 and 7.

Conductivities were measured using an Industrial Instruments Conductivity Bridge Model RC16B2. The conductivity cell (Beckman) used has a cell constant of O.l. Conducting data are also listed in Tables 6 and 7.





(a) Magnesium

(b) Lithium

Figure 6. Oscilloscope Traces of Lithium and Magnesium in $\gamma\textsc{-Butyrolactone}$ (1M KPFs) Electrolyte.

Table 6 FREE ELECTROLYTE CELL DISCHARGE DATA FOR LITHIUM IN NON-QUEOUS ELECTROLYTE

	м	H20	Ga	Gassing		Voltag	Voltage at amp/in. 2 (volts vs NCE)	np/1n.	2 (vol	ts vs	NCE)			CD Cmb	
Electrolyte System	x 10 20 1cm 1	mg/ml	M/o I	IWI	0	.025	.05	0.1	0.2	0.3	4.0	0.5	; >	1n. 2	Notes
1. KPFe ≈ 0.2M AN	1.8	1.5	'n	>>	3.0	2.7	2.3	1.9	6.0				1.1	0.1	White precipitate near
2. Liclo4 im DMF	1.86	2.5	п	ď	3.2	2.6	2.2	5.0	1,2						Gelatinous precipitate
3. Liciim DMF	0.86	1.6	ū	u	3.25	2.95	2.75	2.35	1.2						AD TUT SILICITOR
6. Liclo, im Y-BL	1.14	1.1	E	r	3.1	8.8	2.65	2.25	1.55				9.0	0.1	
9. LICIO4 IM MF	1,22	3.5	¤	и	3.0	2.95	2.90	2.75	2.55	2.40 2.25	2.25	2.15	0.5	0.2	
4. KSGN IM DMF	2.4	0.38	۶		3.25	2.95	2.9	2.8	2.55	2.3	2.0		0.3	0.3	Yellow gelatinous ppt.
5. MPFe 1M DMF	2.65	0.56	ď		3.2	3.15	3.1	3.05	2.95	8.8	2.3		0.32 0.3	6.0	
7. KSCN 1M Y-BL	96.0	49.0	ц		3.25	3.1	3.0	2.8	2.55	₩			0.64 0.2	٥.2	
10. KSCN-0.5M MF	0.16	0.41		≻	3.25	2.85	5.6	2.25	1.33	0.3			1.2	0.2	
11. KPFe IM MF	0.20	0.67	ជ	Ľ	3.25	2.95	5.9	8.8	2.75	2.7	5.6	2.3	0.2	0.2	
8. KPFe 1M Y-BL	0.81	0.55	r	ជ	2.85	2.85 2.55	2. 4.	2.3	2.1	1.95	1.2		4.0	0.2	

Table 7.

PREE ELECTROLYTE CELL DISCHARGE DATA FOR MAGNESIUM IN NONAQUEOUS ELECTROLYTE

	Notes	Improves in use; turns black		-		Grey precipitate	Ppt.	Pat'd etch soln.	Black surface	Black surface spots	Immediate	After Use			0.4 v decay in 4 sec.			
CD amp/	1n. 2		0.05	0.10									0.2	6.0	0.025		,	٥.٥
IR.	>		0.85	0.5									4.0	0.45	7.0		,	0.0
	0.5												0.5	1.30			,	9
VCE)	0.4												9.0	1.44			;	1.22
ts vs 1	0.3								•				0.8	1.60				1.59 1.40
2 (vol	0.2			0.98			0.25						1.1	1.73			i	1.59
mp/1n.	.025 0.05 0.1 0.2 0.3 0.4	0.45	0,45	1,32	2.2		0.91	1.56		0.68	0.48	0.48	1.3	1.89			0.30	1.80
e at	0.05	0.80	1.25	1.5	1. 6		1.05	1.64	1.1	0.75	0.85	1.26	1.5	1.98	0.56		0.50	1.92
Voltag	025	0.95	1.60	1.65	1.8	0.25	1.13	1.82	7.4	0.80	1.26	1.46	1.5	2.02	06.0	0,40	o.5∂	1.98
i	0	1.4	2.2	2.0	5.0	1.05	1.38	1.95		0.75	1.68	2.00	1,62	2,10	1.77	1.16	ુ6 ∙ 0	2.05
	Gassing	No	N _o	*	No	c _N	No	No	Yes	No	No		Yes			No	o _N	Yes
H ₂ 0	mg/m	1.5	1.6	80	1.6	1.1	2.2	1.1	0.38	0.56	0.64		1.7	2.1	0.41	0.67	0.55	9.0
×	x 10 20 cm 1	1.8	0.86	2.6	0.86	1.14	1.22	0.86	2.4	2.65	96.0		3.7	3.1	0.16	0.20	0.81	1.8
		AN	DMF	Меон	DMF	X-BL	MF	DMF	DMF	DMF	(-BL		MeOH	MeoH	MF	MF	-BL	MeOH
Electrolyte	tem	0.2M	И	M12(40	IM	L1ClO4 1M	* JM	П	ЛM	ЛM	μŢ		ML	WI *	Ψī	MI	3M	лм
Elect	Sys	KPF.	L101	Mg(C1(Lici	L1C10,	L1010,	L1C1	KSCN	MPF_{θ}	KSCN		KSCN	L1C10,	KSCN	KPFe 1M	KPFe	LICI
		н	a	13	r	7	10	#	Ŋ	9	80		15	14	11	12		16

* No gassing at open circuit--0.5 ml/min/in.2 (moderate) with current drain.

All solutions were saturated with electrolyte or made one molar when possible. First quarter work with lithium was restricted to free electrolyte testing. A clean lithium surface was cut from a block (Fisher Scientific Co.) and used without further treatment.

c. Discharge of Lithium in Nonaqueous Electrolytes

The results of the free electrolyte half cell tests are shown in Table 6.

The lithium/acetonitrile (AN) system showed considerable gassing although the water content was relatively low. Formation of a white precipitate and poor performance in the presence of KPF_6 electrolyte eliminated this solvent for further work during this quarter.

With N,N-dimethylformamide (DMF), lithium voltages improved. However, a gelatinous precipitate formed on the anode and caused significant IR drop as the discharge progressed. Morpholinium hexafluorophosphate (MPF $_6$) was the best of the solutes tested.

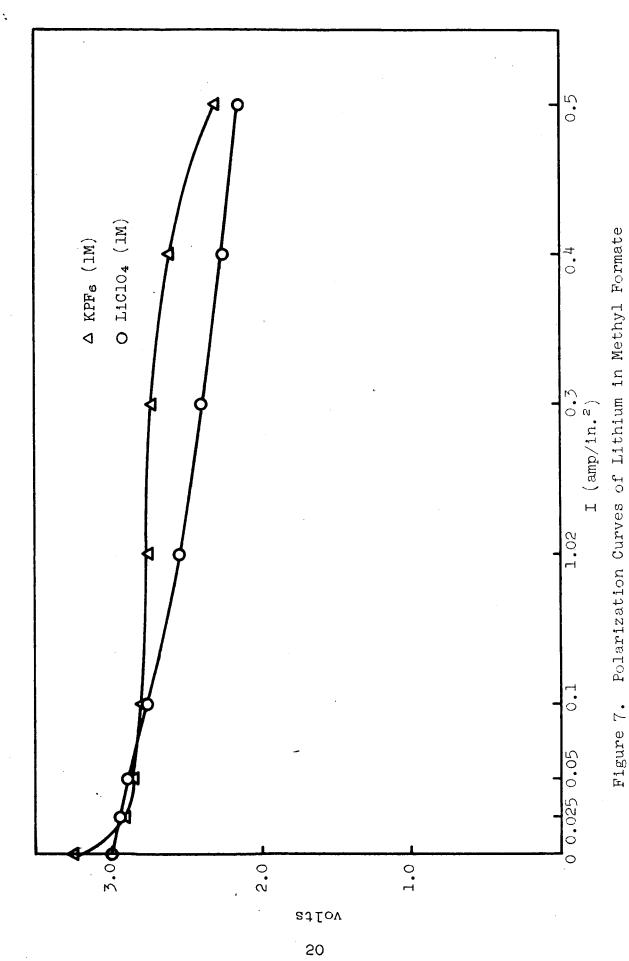
In γ -butyrolactone (γ -BL), the effect of solute (LiClO₄, KSCN, KPF₆) was inconclusive.

Methyl formate with KPF6 or LiClO4 gave the best results with lithium. Lithium performance was somewhat better with KPF6 despite the lower electrolyte conductance. The polarization curves are illustrated in Figure 7. Satisfactory voltages were maintained at current densities up to 0.5 amp/in. 2 .

The Li/LiClO4 lM in methyl formate showed some gassing at open circuit. The open circuit potential was -3.0 v vs NCE. At high currents there was some boiling of the solvent, but true gassing at the lithium did not appear to increase. The IR measurement at 0.2 amp/in.² showed a perfect square wave oscilloscope trace at 10 msec/cm. The IR of 0.5 v added to the operating voltage of 2.55 shows that all the polarization is due to IR. The current was maintained at 0.2 amp/in.² for over 2 hours, with the operating voltage changing randomly between 2.55 and 2.65 v.

With KPFe in the Li/methyl formate systems, potentials were higher and gassing was not observed. However, the lithium surface turned grey. No long-term test has been made with this system.

It is apparent that KSCN decomposes too rapidly to be useful as a solute in any electrolyte system with lithium.



It must be emphasized that these results are from initial screening and are of relative value only. The Luggin capillary was adjusted from $\approx 4 \, \text{mm}$ to $\approx 0.5 \, \text{mm}$ after the following systems were tested: AN (KPF₆), DMF(LiClO₄), DMF (LiCl), γ -BL (LiClO₄).

In general, the oscilloscope traces using lithium anodes showed IR-type polarization only (<1 millisecond to attain final voltage). A typical trace is shown in Figure 6.

2. Magnesium in Nonaqueous Electrolyte

Magnesium in neutral aqueous electrolyte was studied using the free electrolyte cell and tape configuration cell for comparison. The anode material was primary magnesium foil (10 mil) from Dow. The various surface treatments are described individually below.

The OCV of -1.70 v vs NCE for Mg/Mg(ClO₄)₂ aqueous system in the free electrolyte cell test is in agreement with previous static cell data. However, the operating voltage at 0.5 amp/in.² was -1.12 v. An oscilloscope trace showed an IR component of 0.4 v. This operating voltage was less than that of a tape cell and we can, in retrospect, attribute this to lack of cleaning and to lack of chloride ion in solution. Thus, the magnesium may remain somewhat passive. The gassing rate increased under load, and was 1.4 ml/min/in.² at 0.5 amp/in.². The time for complete discharge at this rate was 62 minutes, giving a coulombic efficiency of 86%.

In the tape configuration static test, the silver chloride counter electrode was used.

A capacity of 20 minutes at 0.5 amp/in. was obtained for the AgCl electrode (from the Yardney Silcad® AgO plate) in aqueous Mg(ClO₄)₂ electrolyte versus expanded magnesium. An anode potential of -1.70 v vs NCE was obtained at 0.5 amp/in. drain. This potential includes an IR of 0.15 v as determined by the interrupter circuit (Fig. 11). Using a solid magnesium anode, no appreciable current could be drawn, due apparently to gas blocking. With the same system, but with a gassing platinum screen cathode, much electrolyte had to be added during the test. However, 32 minutes at 0.5 amp/in. was obtained for a solid magnesium anode at an operating potential of -1.5 v vs NCE. This test appears to substantiate the gas blockage theory and also the earlier result that chloride ion improves the magnesium discharge, probably by breaking down the passive magnesium oxide surface film. The magnesium for these anodes was cleaned with detergent or KOH and then chemically polished with 0.1M HCl.

Discharge data for magnesium in nonaqueous electrolyte are difficult to interpret because of sensitivity to surface cleaning methods. In many cases a high open circuit would be

obtained but no appreciable current could be drawn. Electrodes often improved with time under load, or they improved if a higher current was passed initially.

Of the cleaning methods, 3% HCl in DMF was better than 1% HCl in H₂O, which in turn was better than light polishing with emery paper. However, the best method is based on a patented cleaning solution. This solution was used for the free electrolyte cell results in Table 7 except for the AN (KPF₆) and MeOH[Mg(ClO₄)₂] systems.

The etching solution used was:

23 vol-% dioxane

60 vol-% isopropyl alcohol

10 vol-% HNO3 (70% HNO3 in H₂0)

5 vol-% HCl (38% HCl in H₂0)

2 vol-% acetic acid.

The solution was used in the drybox and the magnesium was wiped clean of the etching solution before assembling the cell.

Although the oscilloscope pictures using lithium anodes showed IR-type polarizations only (<1 millisecond to attain final voltage), magnesium anodes showed varied and complicated polarization behavior. The best IR measurement was the initial break from the on-to-off cycle. The off-to-on cycle often showed a larger initial polarization, which may be the IR of the semipassive film on the magnesium. There have been instances in which the trace showed a maximum polarization and then improvement of potential for an off-to-on cycle. Also there have been maxima in the potential of the on-to-off cycle.

Some limited data are available for tape configuration static tests with magnesium in methanol. A Mg/Mg(ClO₄)₂ lM in MeOH system was tested with a gassing platinum cathode and AgCl cathode. The methanol water content was 17 mg/ml and an aqueous HCl etch was used for the magnesium. The data are given in Table 8.

The high water content of the electrolyte and unsatisfactory surface treatment are reflected in the poor anode voltage and high polarization under current drain.

	Pt Catho	de	AgCl Cath	ode
I, amp.	E (v vs NCE)	<u>IR* (v)</u>	E (v vs NCE)	IR (v)
0	-1.48		-1.45	
0.025	-1.47	0.11	-1.34	0.12
0.05	-1.46	0.17	-1.26	0.16
0.1	-1.45	0.23	-1.24	0.32
0.2	-1.43	0.29	-1.13	0.41
0.3	-1.39	0.35	-0.90	0.55
0.4	-1.35	0.39	-0.90	0.68
0.5	-1.20	0.56		

^{*} IR-Anode to calomel on anode side of separator. Obtained with interrupter circuit.

3. Plans for Second Quarter (Anode Research and Development)

The screening of selected lithium and magnesium systems will continue, using the free electrolyte test cell. In addition, this technique will be used to isolate and define problems that may be encountered in full cell tests.

The effect of lithium purity and system water content will be determined.

Development of a lithium anode in a tape configuration will be undertaken. A specific effort will be made to produce a high surface area electrode.

Full cell tests starting with the ACL-85 cathode as counter electrode will be initiated.

III. TASK II - TAPE CELL EVALUATION

A. TAPE CELL PREPARATION

The continuous cathode tape manufacturing machine was serviced and refitted with improved press drying rolls and take-up reels. Production of continuous lengths of ACL-85 tapes will begin early in the second quarter.

B. DYNAMIC TESTING

No dynamic testing was performed in the first quarter. A new test unit was designed and constructed. Complete details are given in Section IV. Dynamic testing of the aqueous Mg/ACL-85 system will begin early in the second quarter.

IV. SUPPORTING RESEARCH

A. TEST EQUIPMENT DESIGN AND IMPROVEMENT

1. Free Electrolyte Half Cell Unit

A special current pulsing switch device has been constructed and is being used with an oscilloscope to measure electrode characteristics during the first few milliseconds of discharge. The switch consists of a mercury-wetted relay that closes the cell current circuit and generates a trigger output to start the oscilloscope.sweep. When the cell current is driven by a properly loaded constant current supply, the oscilloscope display represents the electrode performance at the selected current density for the period of time from open circuit through the first 10 milliseconds of discharge. Figure 8 illustrates the electrical diagram for the system. The test cell itself is shown in Figure 9.

2. Limited Electrolyte Full Cell Static Test Unit

Pulse current equipment has been built and used which is a modification of the Kordesch-Marko bridge and the Allis-Chalmers bridge. The modification used biases the pulse transistor in such a way as to keep it in the cut-off state when high voltage cells (3.0-4.5 volts) are being tested. These modified bridges are capable of measuring the IR-free electrode performance from open circuit to 3-amp discharge current. The Kordesch-Marko bridge records an IR-free voltage reading while the Allis Chalmers bridge records the $V_{\rm IR}$ contribution directly. The circuitry involved is illustrated in Figures 10 and 11.

3. Dynamic Test Station

The new dynamic test station for dry tape batteries has been completed (see Figure 12). The tester is capable of tape speeds of 0.002 to 1.0 in./min. The speed is continuously variable and accurately maintained by a Zero-Max drive system.

An automatic voltage-current recording system has been incorporated to make 24-hour operation possible. The electrolyte feed for the test station is provided by a Milton-Roy controlled volume pump. The electrolyte storage tanks provided have capacity for 24 hours of testing on tape batteries before refilling is required.

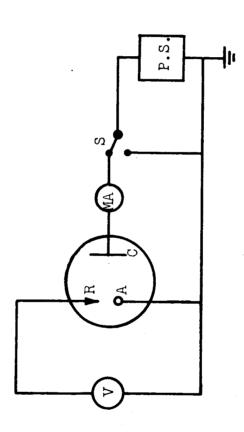
Calibration of the drive system and electrolyte feed system has been completed. Evaluation of the aqueous Mg/ACL-85 tape system will begin early in the second quarter. Compared with the earlier test apparatus, the new dynamic test station has the advantages of long-term unattended operation and more accurate speed and electrolyte feed control.

Tektronix 545-A oscilloscope type L plug in (fast rise calibrated preamp. I meg impedance).

Potter and Bromfield Mercury wetted contact relay JMI 121M. Toggle switch and 90 v dc used to activate relay and trigger CRD. Harrison Laboratories Model 6200A power supply. P.S.

Weston Model 911 milliameter. Platinum auxilliary electrode (cathode).

Test electrode (anode) Reference electrode. A D R A



Circuitry for Free Electrolyte Half Cell Test Figure 8.

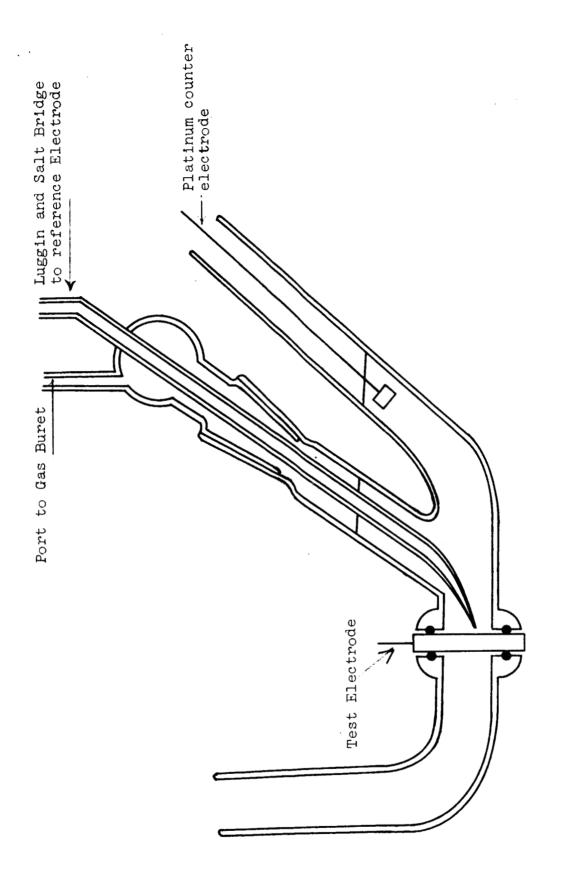
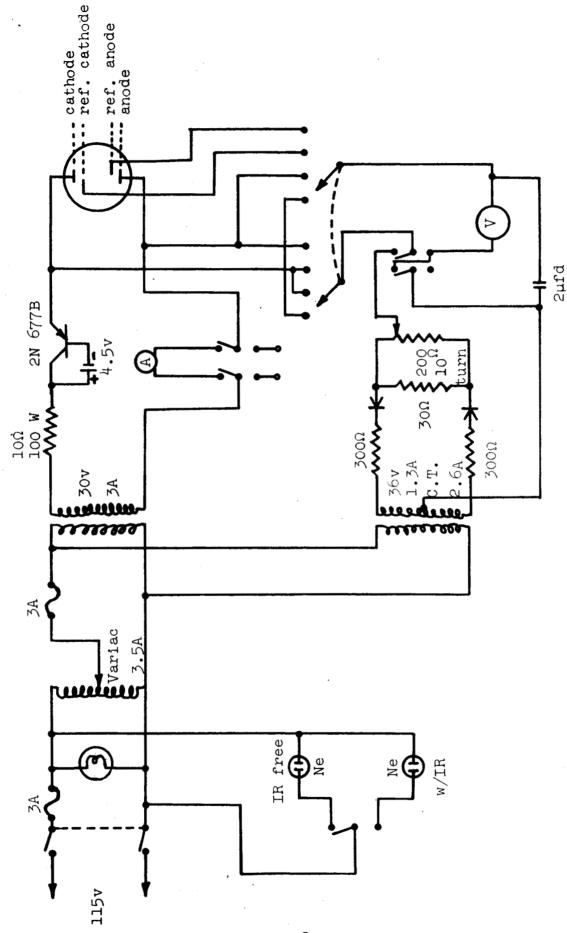
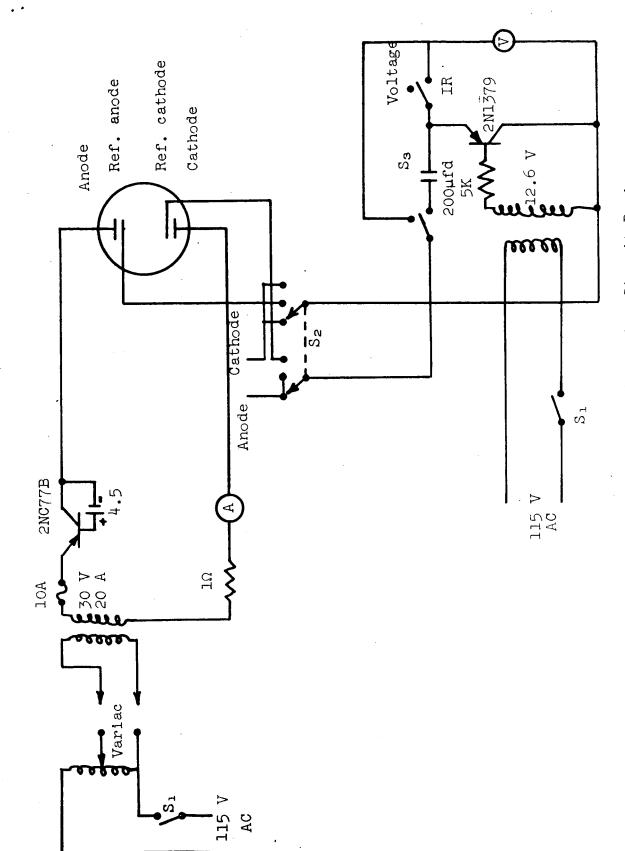


Figure 9. Free Electrolyte Test Cell.



Kordesch-Marko Type Interrupter Circuit Diagram Figure 10.



Allis-Chalmers Type Interrupter Circuit Design Figure 11.



Figure 12. Dynamic Test Unit--Drive, Recorder, and Electrolyte Supply.

4. Improved Water Analysis Techniques

The automatic Karl-Fisher titration currently being used for water analysis has reliability limitations, particularly at very low water concentrations. Several other faster and more peproducible techniques are being investigated with the goal of improving sensitivity. Two techniques examined during this report period were the use of nuclear magnetic resonance and vapor phase chromatography. Nmr was discarded early as a useful technique because its sensitivity limit was approximately 0.1% (too high for our purposes). In addition, its use is restricted to solvents that do not have -OH or N-H bonds.

Two vapor phase chromatographic procedures were investigated:
(a) gas-solid chromatography and (b) gas-liquid chromatography.

a. <u>Gas-Solid Chromatography</u>

Vapor phase chromatograms of water show the phenomenon of tailing, as evidenced by nonsymmetric chromatographic peaks. The high polarity of water accounts for this tailing. The quantitative measurement of chromatographic peak areas is thus subject to tailing-dependent error and the accuracy, and precision of analyses are affected. The possible application of gas-solid chromatography to minimize this error was investigated.

Columns were prepared from Fluoropaks, a perfluorocarbon chromatographic column support (Wilkins Scientific Instruments, Inc.). The column dimensions were 6 ft x 1/4 in. The He carrier gas flow rate was 77 cc/min, and column temperature was constant at 30°C. Injection temperature was 290°C, and the temperature of the hot wire detector was 320°C. Under these operating conditions, a useful separation of water from dimethylformamide does occur and water analyses were possible. Further investigations, however, showed two limitations of this gas-solid chromatographic procedure. First, the procedure was found to be not universal but limited to high boiling solvents (low boiling solvents elute with the water peak and interfere in analyses). Second, gases (e.g., O_2 , N_2) dissolved in the solvent elute just before water and may give interference, particularly at low water concentrations. In view of these limitations and of the desirability of a universally applicable method, the use of gas-solid chromatography was discarded.

b. Gas-Liquid Chromatography

The following column was prepared and investigated for water analysis: 8% Carbowax 20M (Carbide and Carbon Chemicals Co.) on a perfluorocarbon coated diatomaceous earth, Gas-Pack F, 60-80 mesh, (Chemical Research Services, Inc.). Column dimensions were 7 ft x 1/4 in. in 304 stainless steel. Column operating temperature was constant at 100°C and He carrier gas flow rate was

86 cc/min. The instrument was the Perkin-Elmer Vapor Fractometer Model 154. Ten microliters of sample were routinely injected to maximize sensitivity. A typical chromatogram of 0.8% water in dimethylformamide is shown in Figure 13. Retention times are in parentheses for air (0.5 min), unidentified peak, perhaps dimethylamine (1.1 min), water (3.9 min), and dimethylformamide (17.2 min). A series of standards of water in dimethylformamide was prepared and water peak area calibration data were obtained using the method of standard addition.

Peak area measurements are reproducible to \pm 0.05 in. at 0.1% water concentration. This corresponds in this procedure to a precision of \pm 0.01% water at 0.1% water concentration. Some of the preliminary calibration data are shown in Table 9.

The procedure is now being extended to lower water concentrations. The data thus far obtained indicate a limit of detection of 0.001% or 10 ppm water, with an accuracy of \pm 10% of the water concentration.

Preliminary studies of additional solvents have begun. Vapor phase chromatograms of these solvents are shown in Figures 14 (methanol), 15 (methyl formate), 16 (ethyl formate), 17 (y-butyrolactone), and 18 (acetonitrile). The retention times show that this method of water analysis is applicable to these solvents with one limitation. The tailing of acetonitrile, Figure 18, was shown to limit water analysis in acetonitrile to water concentrations above the 1% level. Water in the other five solvents will be detectable at the 0.001% concentration level.

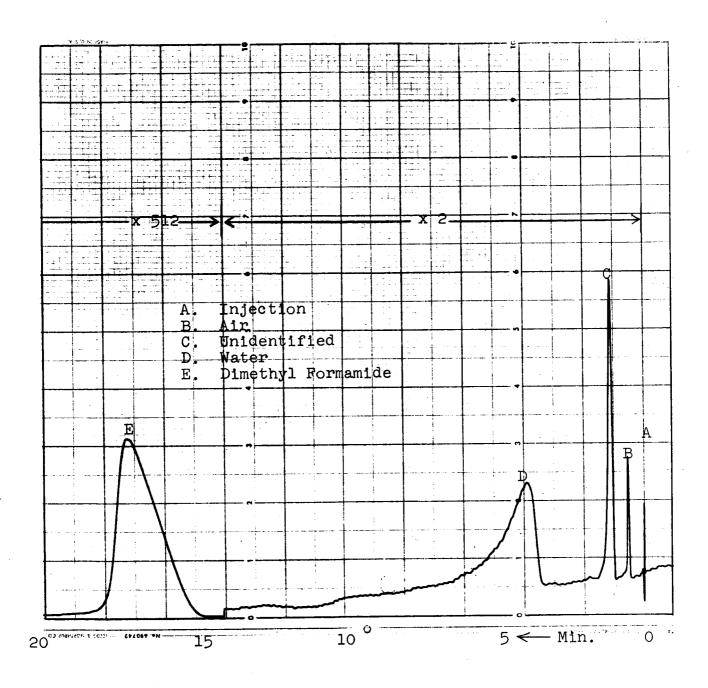


Figure 13. Dimethyl Formamide Vapor Phase Chromatogram.

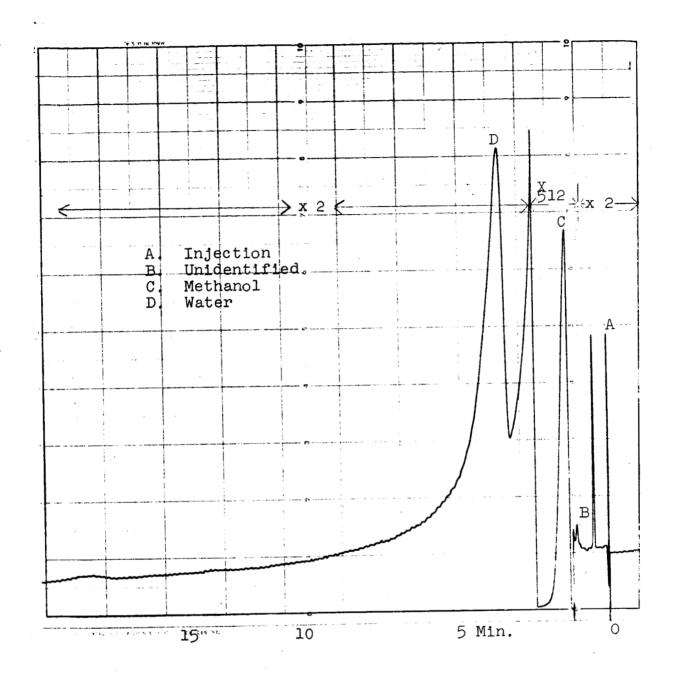


Figure 14. Methanol Chromatogram.

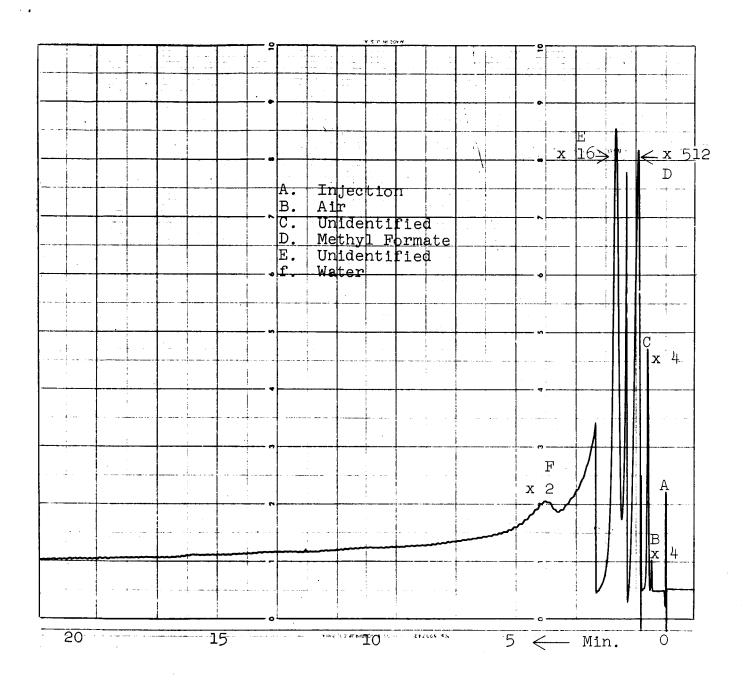


Figure 15. Methyl Formate Chromatogram

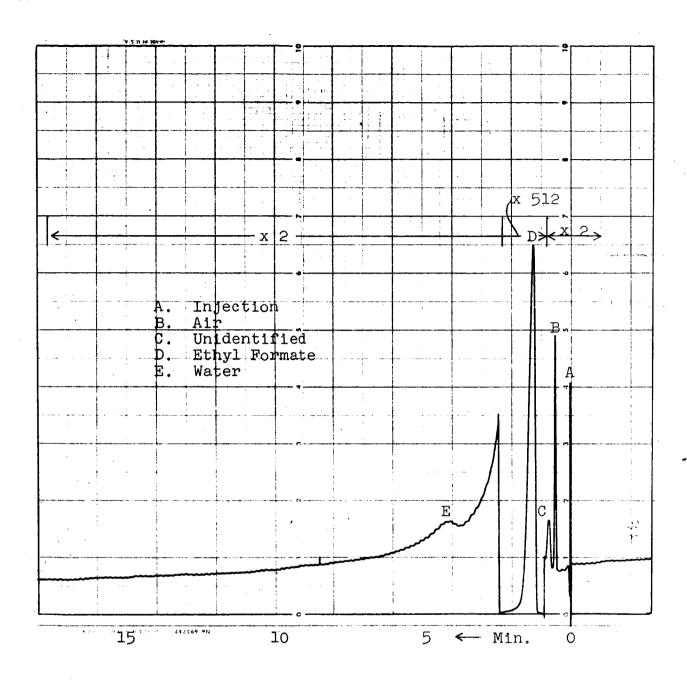


Figure 16. Ethyl Formate Chromatogram.

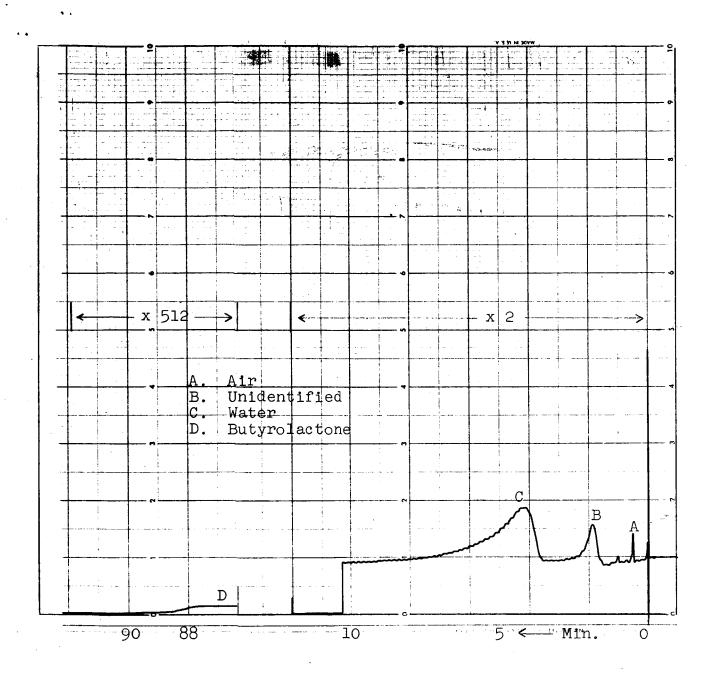


Figure 17. Y-Butyrolactone Chromatogram.

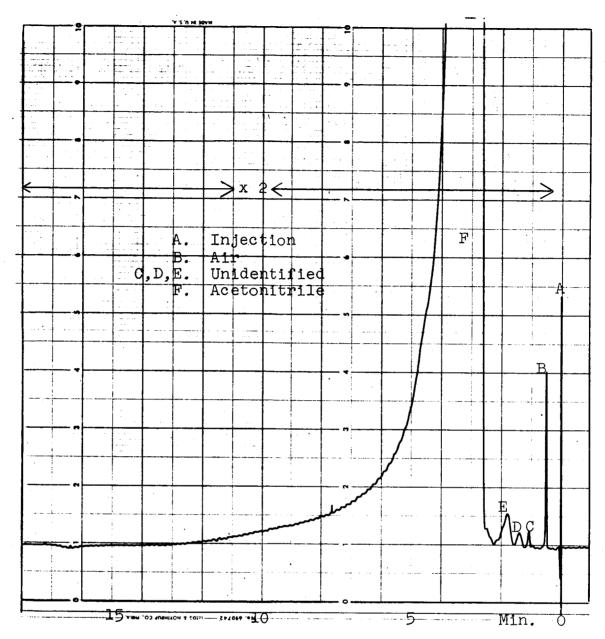


Figure 18. Acetonitrile Chromatogram

Table 9.

PRELIMINARY CALIBRATION DATA FOR VAPOR PHASE CHROMATOGRAPHIC DETERMINATION OF WATER

Peak Area,	Water Concentration,		in.², %
0.398 <u>+</u> 0.045	0.109		3.65
0.582 <u>+</u> 0.026	0.129		4.51
2.043 <u>+</u> 0.020	0.579		<u>3.52</u>
		Mean	3.86 ± 0.40

Table 10.
SOLVENT RETENTION TIMES

Solvent	Retention Time, Min.
Methyl Formate	0.9
Ethyl Formate	1.2
Methanol	1.4
Acetonitrile	3.0
Water	3.9
Dimethylformamide	17.2
γ-Butyrolactone	87.0

V. REFERENCES

1. C. K. Morehouse and R. Glicksman, <u>J. Electrochem. Soc.</u>, <u>104</u>, No. 8, 467 (1957).

Table A-1

STATIC CELL DISCHARGE DATA FOR TRI-N-CHLOROTRIAZINETRIONE TAPE CATHODES

Trichlorotriazinetrione (Monsanto AC1-85®) Acetylene black (Shawinigan)-unless specifically otherwise noted Polyvinyl formal binder (2.75 wt. %) 1/8" dynel fiber (2.75 wt. %) Cathode Composition:

Tape Cell Area = 3 inches x linch

= primary magnesium foll (Dow)

Anode

Separator = 4 M11 Dynel (unless specifically noted)

Remarks		Steady cathode v drop	Steady cathode v drop	=======================================	20% A-625 graph1te	: =	= = : *	Steady cathode emf drop	= =	=======================================	discharge curve	Tape pressed at 1000 ps1			Flat discharge curve		Flat discharge curve	Tape pressed at 1000 ps1
Operating Voltage, v	21.10 1.99 2.09 2.00 3.00	2.10	1.75	 	1.70	2.08	1.72	 26.1	1.00 50.1	1.80	2.45	1.80	1.75	1.70	0, c 0,00	1.70	0. 4. 0. 0.	1.88
Cathode Efficiency.	7. 2. 2. C.	32°E	N N	7.7. 7.7.	/@/ /@/	26 33	19,	52	n c	82	S.	<u>7</u>	N C	× 2	4 K	282	ري د د	24
Current Density Amp/in.2	0000	000	000	, , , , ,	.00	0 C	0.33	0.33	0.00 0.00 0.00	0.00	0.33	0.33	0 5 5 6 6	 	0.33	0.00	0.33	0.5
Electrolyte	2MACl3·1MHCL	2MMgCl2	2MMg(C104)2	r r	2MMgC12	2MMg(C104)2	2MMrC12	2MMg(C104)2	2MMgc12	ZMMBC12 ZMMB(C104)2	2MA1Cl3	2MMgC12	2MMg (C104)2	2MMg(C104)2	2MA IC13	ZIMMEC12 ZIMME(C104)2	2MA1C13	ZMMgC12 ZMMgC12
Theoretical Capacity Amp-Min/in.	8000 6484	nna	เพาะ ได้เ	, w	9.0	٠. د.	, r	0.0	o.		3.0	3.3	4.1 7	, w . r.	4 1 W	∵ ≠	コ! バル(v= 2.ú
Wt.% Active Material	S S S S	32.5	ก็ก้	, <u>, , , , , , , , , , , , , , , , , , </u>	ν. 4. 4.	<u>ئر</u> ي	7.4	<u>.</u>	₹.	ა ქ. ქ	. T.	π	⊅ (วี	000	86	200	301	70
Run No.	78602-6 78602-7 78602-8	78618-4 78618-4	78618-7	78620-7	78624-1	78624-2	78604-4	78623-8	862	78623-10	78623-12	78624-5	78624-6	78627-1	78627-2	78627-5	78627-12	78629-1 78629-2

												.	First vac. dried SAB				Tape stored 24 hrs. in air	: =	E 85	± =	4	after 24	+ + + + + + + + + + + + + + + + + + + +	Tested fresh		=	fresh	ıa '	Tested Iresh	:		Tested after 24 hours	fresh		=	E .	Tested after 24 hours			1221
	O.C.V.	2.62	20.00	8.8 80 80 80 80 80 80 80 80 80 80 80 80 80	0 c	,,	200	0.0	2.60	2.90	2.62	2.92	2.99	2.93	90.00	2.92	2.92	2.91	2.91	20.00	2.97	70	76.	2,07	2.96	2.95	2.95	2.92	9.97	и с И с	0.0	000	2,63	06.2	2.92	2.92	2.96	2,0	, o	2.95
	Operating Voltage,v	1.80	1.85	1.52	1.70	00.0	 8	 		, c		2.30	2.03	2.05	2.15	2.15	2.10	5.00	2.10	1.80	1.80	, o	200	0	2.20	2.05	2.05	2.20	2.25	22.0	07. 01.	0.00	2.15	01.0	1.90	2.00	2.20	2.10	01.70	20.19 190
(Cont.d)	Efficiency	# C	35	31	11	5,6	ر د د	ק ק		\ ²	\ 2	23	57	65	61	59	55	63	2	36	90	200	. 02	- ic	202	54	57	2 π	69	8	2 Q 1 Q	2 6	70	- F	\\ <u>\</u>	29	70	70	20) 99
Table A-1 (Current Density Amp/in.2	0.33	0.53		٠ ان	~ 1	0.175	0.1.0	0.107	79.0	191.0			0.33	0.33	0.33	0.33	0.33	0.33	0.33	0.33	0.0) / / k	0.52	0.33	0.33	0.33	0.33	0.33	0.50	0.00	00.0) , , , , ,		200	0.33	0.50	0.50	0.20
	Electrolyte	2MMg(ClO4)2	ZMA ICIO	2MA1C13	2MMg(C104)2	2MA1C1-	2MAlcla	ZIMING (CIO4/2	ZMMGC12	CM148 (0104 / 2	ZMAIC13	OMA 1 CT & C. SMHCT		=	=	=	=	£	= :	= :	= =	: =	=	=		=	11	=	= :	= :	: :	: =	E	=	ŧ	F	=	= :	= =	: =
	Capacity Amp-Min/in.	ν ν.ι	4 4 M	1 α.	4.5	ر. د	∵ !	ئ	4 = V 6	- -=	v≃ •••) oc	, k	, rc - 00	50.0	ر. د.	, o	4.2	.≠ .∪.	3.1	9.	٠, ن	v.		2 K) K	/4	Γ. 4	3.7	3.7	<u>.</u>	4 t	() = 4 M	^. t	o =	† ¢	-9.	5.3	5.7	9. v.
	Wt.% Active Material																																							20
	Run No.	78629-3	78629-5	78631-1	78631-3	78631-6	78631-7	78631-10	78634-7	78054-8	78654-9	71-66001	78641-7 78651-7	78651	78651-7	- יינעאאַ	78656-6	78656	78656-9	78656-10	78656-14	78659-3	78659-4	78659-5	78561-3	2-T000/	78661-7	78663-10	78665-6	78665-7	78665-4	78665-5	78665-9	78567-5	78667-2	/-/.000/	78667-3	78667-5	78670-10	78670-5 78670-3

1. いいのでは、から

Remarks	A.C. Bridge				Steady cathode emf drop		ady cathode emf d	E E	,	hode emf d	=	= =	= -	Steady cathode emf drop
VS.ME	6.99.99.99.99.99.99.99.99.99.99.99.99.99	000000 000000	70000 2040 401-80	, a a a a a à a à à à à a à a a b a a a	2000 2000 2000 2000 2000	, a a a a y a a a a y a a a a	17.0	22.00	, u u 0,00 0,00 0,00 0,00 0,00 0,00 0,00		2.71	2.91	, a a 0 6 6 0 6 6	2.95 2.95 2.92
Operating Voltage, v	99999999999999999999999999999999999999	ับบบบบ น้ำน้ำ กับบั	ที่ ขั้นเข้า ขั้น	หลดลา ระหากับ ระหากับ	2.1.1.0 0.000 0.000	, a a a a	, , ,	.883	0.0.0	25.20	1.70	2.50	2.10	8.00 8.00 8.00 8.00 8.00 8.00 8.00 8.00
Cathode Efficiency	WWW.	178877 7887 7887	77 77 70 6 70 6 70 6 70 6 70 6 70 6 70 6	87868 800	0 4 4 7 0 1 8 0	2500	/ () = // () =	1 t d 1000 1 t	64 64 70 70	-80 -20 -20 -20 -20 -20 -20 -20 -20 -20 -2	ታ የ ያ	, E.	かんしょ	750 750 750
Current Density Amp/in.2	00000	0000 20000 20000	0.00 .50 .57 .00 .00 .00 .00 .00 .00 .00 .00 .00 .0	00001 00077 0001			00.00	000 04.0 04.0	101 0.40 101 101				e)	Micla 0.67
Electrolyte		: : : :	::::		2MMgCl2 2MMgCl2	2MAlcl ₃ ·0.5MHCl "	2MMgC12	2MM&Cla 2MM&Cla 2MM&Cla	2MAIC13.0.5MHC1	IMAICI3 IMMECIZ	1MA1C13·1MMgC	IMAICIS-IMMg(CIO4)2	1.3MMg(C104 b.0.7MA1Cla 1.3MMgC1z.0.7MA1Cla 2MA1Cl	ZMAICIS ZMAICIS 1.3MMg(C104/2.0.7MAICIS 1.3MMgC1z.0.7MAICIS
Theoretical Capacity Amp-Min/in.2	ಇಇ ಬળ≀ ಇಹಬಹು	ບຸ ຈ ເບເບ⇒ ນ່ວໝໍທີ່ເບ	w≠ w∞ ir∞ i o	←≒≒ห,น ด์เข่ต์เข่∞	inua iraio	⊬4 N/Q Ô'N'À'N'	գ տ Գ ւ	すらす	(U.T.)	v.v.	W.	+ 1~	01 Q 1	-000 F
Wt.%Active Material	L			0027										
Run No.	78670-7 78670-9 78670-4 78670-4	78673-8 78673-6 78673-10 78673-10	78673-2 78673-3 78673-1	78677-5 78677-9 78677-3 78677-2	78677-4 78677-8 78677-8	78678-6 78678-1 78678-1 78678-5	78678-2 78678-3	78678-4 78678-9 78685-7	78685-5 78685-10	78685-3 78685-4	78685-1	78585-3 78685-2	78688-6 78688-2	78688-1 78688-1 78688-1

Remarks	Steeds outhods onf dron	מינה שהטווטפט להפסט ב				Stored five days in air	= = =							44.00	Scored chiree days the art.	1000	Stored one day in air										
O.C.V.	2.93 2.93	2.95	0.0 0.0	26.0	2.93	2.95	, c	26.90	2.91	2.95	2.91	2.93	2.94	9.00	9.0		26.5	9.0	26.00	v (, v	20.0	26.2	9.0 9.0	20.0	0 0 0 0	7.75
Operating Voltage,v	9.00 4.00 50.00	1.95	2.35	, v	2.35	2.40	 で	7.0	25.5	2.35	2°.3	2.40	0.40	00.00 00.00	2.20	2.50	2.25	2.20	2.25	8.50	02.20	2.25	2.50	2.30	2.15	2.10	CT • 2
Cathode Efficiency	76 69 69	38	<u> 2</u> 9	57 7.4	7.2	ر ن عرب	χ. Φ.	0 t	ેં	90	63	57,	2.4	, Š	χ, Σ	<u>ڻ</u>	61	62	20	8,	8,	66	وج	64	46	i Si	č
Current Density Electrolyte Amp/in.	2MAICLS .O. SMHCI 0.67		_	79.0 "				-	2MAIC13.0.1MHC1 0.50	2.0 CMA1C1 2.0.25MHC1 0.50		2MA1C13.0.5MHC1 0.50		2MAICIS.O.1MHC1 0.67				.0.2MHC1		0.25MHC1		2MAlcla.0.1MHCl 0.50		2MA1Claro.5MHCl 0.50			2MA1C13 0.11MHC1 0.67
Theoretical Capacity Amp-Min/in.	7.7.	~. 	10.0	n n	140	9.9	10.0	9.5	o o	να) c	o ∞	0.8	8.0	8.0	9.1	9.1	2.6	11.1	8.3	8.7	4.9	9.1	9.5	11.0	11.2	10.0
Wt.%Active Material	65 65 65	o o o	65	č,	O G	90	65	. 65	ري اري	o d O d	C u	9.0 J.C	20	65	65	70	20.	20	70	20	202	20.	70	202	20	70	20
nn No.	8688-3 8690-6	8690-5 8690-4	8690-3	8690-1	8690-8	8690-7	8690-9	8690-10	8693-9	8693-10 8603-1	000 000 000 000 000 000 000 000 000 00	8693-1	01-2098	8693-11	8693-6	8694-9	9-1-098	8694-11	8694-12	8694-8	2-4-698	8694-2	18694-1	8694-5	8694-10	8694-3	4-4698

Table A-2.

ANALYTICAL METHOD FOR THE DETERMINATION OF AVAILABLE CHLORINE

The determination of available chlorine in formulated products containing chlorinated cyanuric acid is performed by means of an iodometric titration with sodium thiosulfate to a starch end-point.

The method given below is recommended.

1. Reagents and Equipment

O.lN standard sodium thiosulfate solution

3% potassium iodide solution
1:1 sulfuric acid
Starch indicator solution
50-ml burette
500-ml Erlenmeyer flask
250-ml graduate cylinder
25-ml graduate cylinder
Magnetic stirrer
Teflon-covered stirring bar
Distilled water wash-bottle
Glassine weighing paper
Analytical balance

2. Procedure

Using the analytical balance, weigh onto a glassine weighing paper the calculated amount of sample for an approximate 40-ml sodium thiosulfate titration. Transfer the sample to a 500-ml Erlenmeyer flask containing 250 ml of 3% potassium iodide solution. Rinse the weighing paper and the walls of the flask with distilled water. Add a stirring bar covered with DuPont's "Teflon" to the flask and place it on a magnetic stirrer. Continue to stir until solution has been effected (usually 3-5 minutes). Add approximately 50 g of crushed ice, and then add 25 ml of 1:1 sulfuric acid. (Caution: when the formulation contains a carbonate, the acid should be added with extra care in order to avoid an excessive rate of CO2 evolution.) Titrate with O.1N standard sodium thiosulfate to a light yellow color while continuing to stir the contents of the flask. Add starch indicator solution to give a deep purple color, and continue to titrate slowly until the color just disappears. Record the volume of sodium thiosulfate solution used, and calculate the available chlorine level according to the following formula:

% Available Chlorine = ml_thiosulfate x Nthiosulfate x 0.03546 x 100 wt sample ACL-85